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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
09/349,713	07/08/1999	HERWIG BUCHHOLZ	MERCK-1900	7039
23599	7590 05/19/2004		EXAMINER	
MILLEN, WHITE, ZELANO & BRANIGAN, P.C.			SHARAREH, SHAHNAM J	
2200 CLARI SUITE 1400	ENDON BLVD.		ART UNIT	PAPER NUMBER
ARLINGTON, VA 22201			1617	

DATE MAILED: 05/19/2004

Please find below and/or attached an Office communication concerning this application or proceeding.

		Application No.	Applicant(s)			
Office Action Summary		09/349,713	BUCHHOLZ ET AL.			
		Examiner	Art Unit			
		Shahnam Sharareh	1617			
The MAILING Period for Reply	DATE of this communication app	ears on the cover sheet with the c	orrespondence address			
THE MAILING DATI - Extensions of time may be after SIX (6) MONTHS fro - If the period for reply spec - If NO period for reply is sy - Failure to reply within the Any reply received by the	ATUTORY PERIOD FOR REPLY E OF THIS COMMUNICATION. e available under the provisions of 37 CFR 1.13 m the mailing date of this communication: iffied above is less than thirty (30) days, a reply secified above, the maximum statutory period w set or extended period for reply will, by statute, Office later than three months after the mailing ment. See 37 CFR 1.704(b).	6(a). In no event, however, may a reply be tin within the statutory minimum of thirty (30) day ill apply and will expire SIX (6) MONTHS from cause the application to become ABANDONE	nely filed s will be considered timely. the mailing date of this communication. D (35 U.S.C. § 133).			
Status						
1) Responsive to	Responsive to communication(s) filed on <u>7/2003, 1/2004</u> .					
2a) ☐ This action is	This action is FINAL . 2b)⊠ This action is non-final.					
	☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is					
closed in acco	closed in accordance with the practice under Ex parte Quayle, 1935 C.D. 11, 453 O.G. 213.					
Disposition of Claims						
4) Claim(s) <u>1-10</u> ,	4) Claim(s) 1-10,12-14 and 26 is/are pending in the application.					
	4a) Of the above claim(s) is/are withdrawn from consideration.					
<u> </u>	Claim(s) is/are allowed.					
	Claim(s) <u>1-10,12-14 and 26</u> is/are rejected.					
7) ☐ Claim(s)	_ is/are objected to. _ are subject to restriction and/or	election requirement				
	_ are subject to restriction and/or	election requirement.				
Application Papers						
9) The specification is objected to by the Examiner.						
10) The drawing(s) filed on is/are: a) accepted or b) objected to by the Examiner.						
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a). Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).						
	claration is objected to by the Ex					
Priority under 35 U.S.C	C. § 119					
	ent is made of a claim for foreign	priority under 35 U.S.C. § 119(a))-(d) or (f).			
a) ☐ All b) ☐ Some * c) ☐ None of: 1. ☐ Certified copies of the priority documents have been received.						
	copies of the priority documents		on No.			
	of the certified copies of the prior					
applicat	ion from the International Bureau	(PCT Rule 17.2(a)).				
* See the attache	d detailed Office action for a list of	of the certified copies not receive	ed.			
Attachment(s)	(, 				
 Notice of References C D Notice of Draftsperson's 	ited (PTO-892) s Patent Drawing Review (PTO-948)	4) 🔲 Interview Summary Paper No(s)/Mail Da				
	Statement(s) (PTO-1449 or PTO/SB/08)		atent Application (PTO-152)			

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Continued Examination Under 37 CFR 1.114

A request for continued examination under 37 CFR 1.114, including the fee set forth in 37 CFR 1.17(e), was filed in this application after final rejection. Since this application is eligible for continued examination under 37 CFR 1.114, and the fee set forth in 37 CFR 1.17(e) has been timely paid, the finality of the previous Office action has been withdrawn pursuant to 37 CFR 1.114. Applicant's submission filed on January 27, 2004 has been entered.

Amendment filed on June 1, 2003 has been entered. Claims 1-10, 12-14, 26 are pending. Any rejection that is not addressed here is considered obviated in view of the amendments.

Claim Objections

Claims 1 and all dependent claims thereof are objected to because of the following minor informalities: the limitation of "a glycoside of any of the above other components," is confusing. It is not clear to which other components is applicant referring. Appropriate correction is required to clarify this language.

Double Patenting

The nonstatutory double patenting rejection is based on a judicially created doctrine grounded in public policy (a policy reflected in the statute) so as to prevent the unjustified or improper timewise extension of the "right to exclude" granted by a patent and to prevent possible harassment by multiple assignees. See *In re Goodman*, 11 F.3d 1046, 29 USPQ2d 2010 (Fed. Cir. 1993); *In re Longi*, 759 F.2d 887, 225 USPQ 645 (Fed. Cir. 1985); *In re Van Ornum*, 686 F.2d 937, 214 USPQ 761 (CCPA 1982); *In re Vogel*, 422 F.2d 438, 164 USPQ 619 (CCPA 1970);and, *In re Thorington*, 418 F.2d 528, 163 USPQ 644 (CCPA 1969).

A timely filed terminal disclaimer in compliance with 37 CFR 1.321(c) may be used to overcome an actual or provisional rejection based on a nonstatutory double patenting ground provided the conflicting application or patent is shown to be commonly owned with this application. See 37 CFR 1.130(b).

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Effective January 1, 1994, a registered attorney or agent of record may sign a terminal disclaimer. A terminal disclaimer signed by the assignee must fully comply with 37 CFR 3.73(b).

Claims 1-10, 12-14, 26 are rejected under the judicially created doctrine of obviousness-type double patenting as being unpatentable over claim1-24 of U.S. Patent No. 6,491,948. Although the conflicting claims are not identical, they are not patentably distinct from each other because the scope of pending claims are anticipated with those already patented.

Claim 1 encompass pharmaceutical compositions comprising isoquercitrin and ascorbic acid. The patented claims are directed to oral formulations of isoquercitrin and ascorbic acid. Accordingly, one of ordinary skill in the art in possession of the patented claims would have practiced the instant pending claims.

Claim Rejections - 35 USC § 103

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.

The factual inquiries set forth in *Graham* v. *John Deere Co.*, 383 U.S. 1, 148 USPQ 459 (1966), that are applied for establishing a background for determining obviousness under 35 U.S.C. 103(a) are summarized as follows:

- 1. Determining the scope and contents of the prior art.
- 2. Ascertaining the differences between the prior art and the claims at issue.
- 3. Resolving the level of ordinary skill in the pertinent art.
- 4. Considering objective evidence present in the application indicating obviousness or nonobviousness.

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This application currently names joint inventors. In considering patentability of the claims under 35 U.S.C. 103(a), the examiner presumes that the subject matter of the various claims was commonly owned at the time any inventions covered therein were made absent any evidence to the contrary. Applicant is advised of the obligation under 37 CFR 1.56 to point out the inventor and invention dates of each claim that was not commonly owned at the time a later invention was made in order for the examiner to consider the applicability of 35 U.S.C. 103(c) and potential 35 U.S.C. 102(e), (f) or (g) prior art under 35 U.S.C. 103(a).

Claims 1-4, 6-8, 12, 26 are rejected under 35 U.S.C. 103(a) as being unpatentable over Abou-Karam et al (J Nat Prod. 1992, Oct; 55(10):1525-7) in view of Hovi et al (Antiviral Research 27 (1995) 263-270) and Bean US Patent 4,132,782.

Abou-Karam teaches that isoquercitrin has profound activity against simplex virus. Abou-Karam also teaches suitable antiviral doses of isoquercitrin (see abstract, and page 1526). Abou-Karam doesn't teach the use of isoquercitrin with ascorbic acid.

Hovi teaches that ascorbic acid has substantial antiviral effects against herpes simplex virus (abstract, pages 268-270).

Bean is further used to show that isoquercitrin containing topical compositions with carotene and other vitamins are known in the art to be useful for suppressing herpes simplex virus.

It is *prima facie* obvious to combine two compositions each of which is taught by prior art to be useful for same purpose in order to form third composition that is to be used for very same purpose. *In re Kerkhoven*, 205 USPQ 1069 (CCPA 1980). Claims

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that require no more than mixing together of two conventional compounds used for the same purpose would be *prima facia* obvious.

In the instant case, it would have been obvious to one of ordinary skill in the art at the time of invention to combine isoquercitrin of Aou-Karam at suitable anti herpes virus activity with ascorbic acid of Hovi, because mixing these two elements to create a third composition for treatment of herpes simplex virus would have been well within purview of an ordinary artisan and further as taught by Bean, the ordinary skill in the art would have had a reasonable expectation of success in using a topical formulation containing isoquercitrin for treatment of herpes simplex virus.

Claims 1-10, 12-14, 26 are rejected under 35 U.S.C. 103(a) as being unpatentable over Abou-Karam et al (J Nat Prod. 1992, Oct;55(10):1525-7) in view of Hovi et al (Antiviral Research 27 (1995) 263-270) and Bean US Patent 4,132,782 as applied to claims 1-4, 6-8, 12, 26 above, and further in view of Lanzendorfer et al WO 96/18381.

The combined teachings of Abou-Karam, Hovi and Bean are described above.

The combined teachings of the cited references do not explicitly teach oral formulation of isoquercitrin or topical formulations comprising a UV filtering agent.

Lanzendorfer is used to show that conventional nature of art in combining flavones in oral and topical compositions. Lanzendorfer discloses topical and oral flavonoid containing compositions comprising various vitamins, UVB or UVA filters and at least one flavone including isoquecertin derivatives. Given the fact that isoquecertin is one of the four subspecies of quecertin, the teachings of Lanzendorfer also encompass

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utility of isoquecertin. Lanzendorfer also indicates antiviral activity with his composition (see abstract, page 3-5, 43, 54-60, claims 1-7, and examples, example 27 is an oral preparation). Lanzendorfer does not specifically teach combining isoquercitrin with ascorbic acid, carotene or other vitamins as an antiviral formulation.

Nevertheless, it would have been obvious to one of ordinary skill in the art at the time of invention to use the teachings of Lanzendorfer and modify the combination of isoquercitrin and ascorbic acid of Abou-Karma and Hovi for the purposes of preparing a topical composition of isoquercitrin with a UV filter, because as taught by Lanzendorfer preparing such topical formulations of flavones well within purview of one of ordinary skill in the art, and one of ordinary skill in the art would have been motivated to formulate topical formulations to reduce systemic side effects of antivirus drugs.

Further, it would have been obvious to one of ordinary skill in the art at the time of invention to prepare such compositions in oral form because formulating such compositions with a new carrier system is a matter of design choice and well within the level of an ordinary artisan.

Conclusion

No claims are allowed.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Shahnam Sharareh whose telephone number is 571-272-0630. The examiner can normally be reached on 8:30 am - 6:00 pm.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Sreenivasan Padmanabhan, PhD can be reached on 571-272-0629. The

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fax phone number for the organization where this application or proceeding is assigned is 703-872-9306.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

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